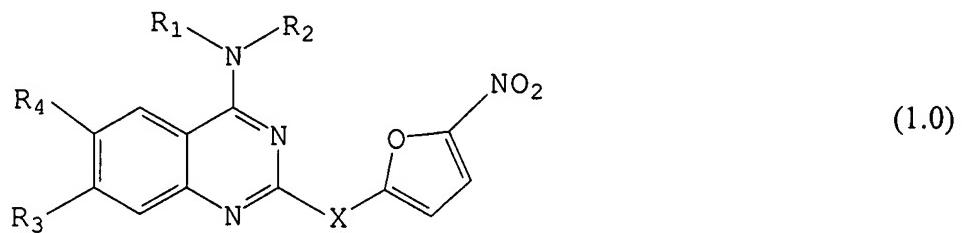


**LISTING OF CLAIMS**

The following listing of claims will replace all prior versions, and listings of claims in the application:

1. **(Original)** A compound of the formula



wherein

X is absent or trans or cis CHCH,

R<sub>1</sub> is (C<sub>1</sub>-C<sub>10</sub>) alkyl unsubstituted or substituted by one to three hydroxy, (C<sub>1</sub>-C<sub>10</sub>) alkenyl unsubstituted or substituted by one to three hydroxy, (C<sub>1</sub>-C<sub>10</sub>) alkynyl unsubstituted or substituted by one to three hydroxy, or aryl unsubstituted or substituted by one to three hydroxy;

R<sub>2</sub> is hydrogen, alkyl or aryl;

R<sub>3</sub> and R<sub>4</sub> are, independently of each other, H, halogen, or a solubilizing group,

with the proviso that at least one of R<sub>3</sub> and R<sub>4</sub> is halogen;

or a pharmaceutically acceptable salt thereof.

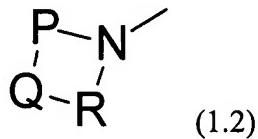
2. **(Currently amended)** [[A]]The compound according to claim 1, wherein R<sub>1</sub> is aryl unsubstituted or substituted by one to three hydroxy and R<sub>2</sub> is hydrogen.

3. **(Canceled).**

4. **(Currently amended)** [[A]]The compound according to ~~any one of claims 1 to 3~~claim 1, wherein R<sub>4</sub> is a halogen.

5. (Canceled).

6. (Currently amended) [[A]]The compound according to ~~any one of claims 1 to 3~~claim 1, wherein the solubilizing group of R<sub>3</sub> or R<sub>4</sub> is



wherein:

P and R are each independently selected from CH<sub>2</sub>, CH<sub>2</sub>CH<sub>2</sub> and CH<sub>2</sub>CHT where T is alkyl, and

Q is O, S, NH or NCH<sub>3</sub>.

7. (Currently amended) [[A]]The compound according to claim 6, wherein R<sub>3</sub> is a halogen and R<sub>4</sub> is partial formula (1.2) wherein Q is NH or NCH<sub>3</sub>.

8. (Canceled).

9. (Currently amended) [[A]]The compound according to ~~any one of claims 1 to 8~~claim 1, wherein R<sub>3</sub> is an amine containing heterocycle.

10. (Currently amended) [[A]]The compound according to ~~any one of claims 1 to 8~~claim 1, wherein R<sub>3</sub> is N-methylpiperazine.

11. (Currently amended) [[A]]The compound according to ~~any one of claims 1 to 10~~claim 1, wherein X is trans CHCH.

12. (Currently amended) [[A]]The compound according to ~~any one of claims 1 to 11~~claim 1, wherein R<sub>1</sub> is hydroxyethanol, hydroxyaniline, hydroxyphenyl, 2-hydroxyethanol, 4-hydroxyaniline, or 4-hydroxyphenyl.

13-17. (Canceled).

18. (Currently amended) [[A]]The compound according to ~~any one of claims 1 to 17~~claim 1, wherein R<sub>2</sub> is phenyl, substituted phenyl, pyranyl, substituted pyridinyl, thiophenyl, substituted

thiophenyl, furanyl, substituted furanyl, thiazole, oxazole or substituted or unsubstituted imidazole.

19. (Currently amended) [[A]]The compound according to claim 12 or claim 15, wherein R<sub>2</sub> is N-alkyl imidazole.

20. (Currently amended) [[A]]The compound according to claim 1 of the formula 6-fluoro-2-[2-(5-nitro-2-furyl) vinyl]-4-(p-hydroxyanilino)-quinazoline.

21. (Currently amended) [[A]]The compound according to claim 1 of the formula 7-(4-methylpiperazino)-6-fluoro-2-[2-(5-nitro-2-furyl) vinyl]-4-(p-hydroxyanilino)-quinazoline.

22. (Currently amended) [[A]]The compound according to claim 1 of the formula 6-fluoro-2-[2-(5-nitro-2-furyl) vinyl]-4-chloroquinazoline.

23. (Currently amended) [[A]]The compound according to claim 1 of the formula 7-(4-methyl piperazino)-6-fluoro-2-[2-(5-nitro-2-furyl) vinyl]-4-chloroquinazoline.

24. (Currently amended) [[A]]The compound according to claim 1 of the formula 6-fluoro-2-[2-(5-nitro-2-furyl) vinyl]-4-(3H)quinazolinone.

25. (Currently amended) [[A]]The compound according to claim 1 of the formula 7-(4-methylpiperazino)-6-fluoro-2-[2-(5-nitro-2-furyl) vinyl]-4-(3H) quinazolinone.

26-28. (Canceled).

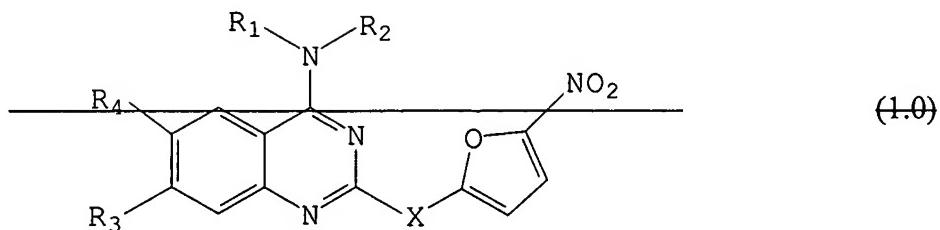
29. (Currently amended) A method for preventing or treating a bacterial infection in a human or an animal, comprising administering to said human or said animal a prophylactically or therapeutically effective amount of a compound according to any one of claims 1 to 24claim 1, effective in preventing or treating the bacterial infection.

30. (Canceled).

31. (Currently amended) A method for antisepsis of, or disinfecting or sterilizing the surface of an object, including a human, of bacteria, comprising: contacting the object with the compound according to any one of claims 1 to 24claim 1 in an amount and for a time sufficient to achieve a desired degree of antisepsis, disinfection or sterilization.

32-38. (Canceled).

39. (Currently amended) A process for the preparation of the[[a]] compound according to claim 1, of formula 1.0



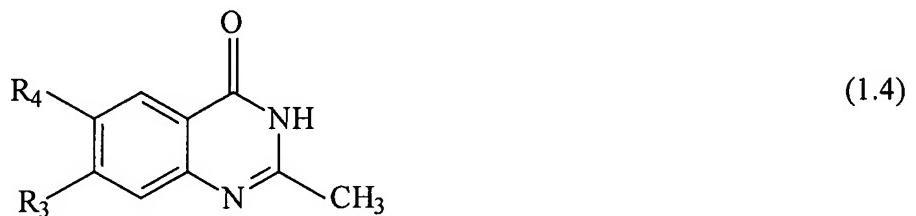
wherein R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are as defined in claim 1,

the process comprising:

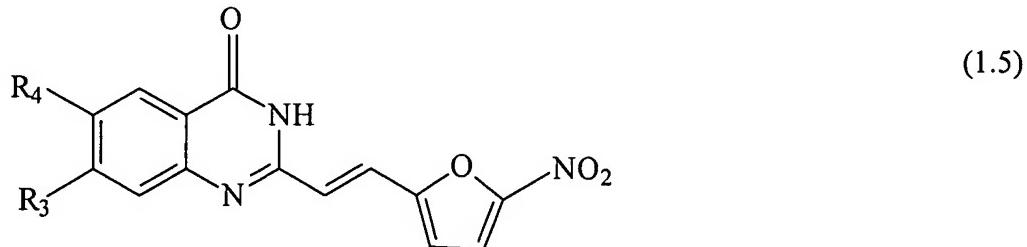
a) optionally reacting a compound of formula (1. 3)



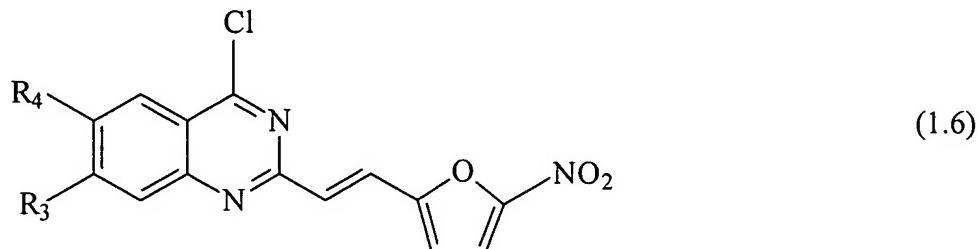
with hydrochloric acid, acetic anhydride and aqueous ammonia, to form a compound of formula (1.4)



b) optionally reacting the compound of formula 1.4 with 5-nitro-2-furancarboxaldehyde, to form a compound of formula (1.5)



c) optionally reacting the compound of formula 1.5 with phosphorus pentachloride and phosphorus oxychloride to form a compound of formula (1.6)



and

d) reacting the compound of formula 1.6 with a compound of the formula (1.7)



wherein X is H and R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, and R<sub>4</sub> are as defined above in claim 1.

40-42. (Cancelled).